

03/31/98

UTILITY PATENT APPLICATION TRANSMITTAL

(Only for new nonprovisional applications under 37 C.F.R. 1.53(b))

Attorney Docket No. 0151/00211 <i>First Named Inventor or Application Identifier</i> Johannsson <i>Express Mail Label No.</i>	
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APPLICATION ELEMENTS

See MPEP chapter 600 concerning utility patent application contents

Assistant Commissioner for Patents
ADDRESS TO: Box Applications
Washington, DC 20231

1. <input checked="" type="checkbox"/> Fee Transmittal Form <i>(Submit an original, and a duplicate for fee processing)</i> 2. <input checked="" type="checkbox"/> Specification [Total Pages [27]] <i>(preferred arrangement set forth below)</i> <ul style="list-style-type: none"> - Descriptive title of the invention - Cross References to Related Applications - Statement Regarding Fed sponsored R & D - Reference to Microfiche Appendix - Background of the Invention - Brief Summary of the Invention - Brief Description of the Drawings <i>(if filed)</i> - Detailed Description - Claim(s) - Abstract of the Disclosure 3. <input type="checkbox"/> Drawing(s) <i>(35 USC 113)</i> [Total Pages []] 4. Oath or Declaration [Total Pages []] <ul style="list-style-type: none"> a. <input type="checkbox"/> Newly executed (original or copy) b. <input type="checkbox"/> Copy from a prior application (37 CFR 1.63(d)) <i>(for continuation/divisional with Box 17 completed)</i> <ul style="list-style-type: none"> <input type="checkbox"/> DELETION OF INVENTOR(S) Signed statement attached deleting inventor(s) named in the prior application, see 37 CFR 1.63(d)(2) and 1.33(b) 5. <input type="checkbox"/> Incorporation By Reference <i>/useable if Box 4b is checked/</i> The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under Box 4b, is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein	6. <input type="checkbox"/> Microfiche Computer Program <i>(Appendix)</i> 7. Nucleotide and/or Amino Acid Sequence Submission <i>(if applicable, all necessary)</i> <ul style="list-style-type: none"> a. <input type="checkbox"/> Computer Readable Copy b. <input type="checkbox"/> Paper Copy <i>(identical to computer copy)</i> c. <input type="checkbox"/> Statement verifying identity of above copies
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ACCOMPANYING APPLICATION PARTS

8. Assignment Papers (cover sheet & document(s))
9. 37 CFR 3.73(b) Statement Power of Attorney
10. English Translation Document *(if applicable)*
11. Information Disclosure Statement (IDS)/PTO-1449 Copies of IDS Citations
12. Preliminary Amendment
13. Return Receipt Postcard (MPEP 503)
(Should be specifically itemized)
14. Small Entity Statement filed in prior application, Statement(s) Status still proper and desired
15. Certified Copy of Priority Document(s)
(if foreign priority is claimed)
16. Other:

17. If a CONTINUING APPLICATION, check appropriate box and supply the requisite information:

 Continuation Divisional Continuation-in-part (CIP) of prior application No. 08/857,245
18. CORRESPONDENCE ADDRESS

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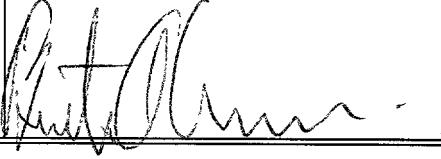
FEE TRANSMITTAL

		Complete If Known	
		Application Number	
		Filing Date	
		First Named Inventor	Johannsson
		Group Art Unit	
		Examiner Name	
TOTAL AMOUNT OF PAYMENT	(\\$)770	Attorney Docket Number	0151/00211

METHOD OF PAYMENT (check one)		FEE CALCULATION (continued)			
1 <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge indicated fee and credit any over payments to Deposit Account Number		22-0185			
Deposit Account Name		Pollock, Vande Sande & Priddy			
<input checked="" type="checkbox"/> Charge Any Additional Fee Required Under 37 CFR 1.16 and 1.17 <input type="checkbox"/> Charge the Issue Fee Set in 37 CFR 1.18 at the Filing of the Notice of Allowance 37 CFR 1.311(b)					
2 <input checked="" type="checkbox"/> Payment Enclosed <input checked="" type="checkbox"/> Check <input type="checkbox"/> Money Order <input type="checkbox"/> Other					
FEE CALCULATION (fees effective 10/01/97)					
1. FILING FEE					
Large Fee Code	Entity Fee	Small Entity Fee Code	Entity Fee	Fee Description	Fee Paid
101	790	201	395	Utility filing fee	\$790
106	330	206	165	Design filing fee	
107	540	207	270	Plant filing fee	
108	790	208	395	Reissue filing fee	
114	150	214	75	Provisional filing fee	
SUBTOTAL (1)			\$790		
2. CLAIMS					
Extra claims	below	Fee from Fee Paid			
Total Claims (18) - 20 =		[0]	X	[\\$22]	= [\\$0]
Independent (2) - 3 =		[0]	X	[\\$82]	= [\\$0]
Claims					
Multiple Dependent Claims	[]	X	[]	= []	
Large Entity Fee Code (\$)					
Small Entity Fee Code (\$)					
103	22	203	11	Claims in excess of 20	
102	82	202	41	Independent claims in excess of 3	
104	270	204	135	Multiple dependent claim	
109	82	209	41	Reissue independent claims over original patent	
110	22	210	11	Reissue claims in excess of 20 and over original patent	
SUBTOTAL (2) [\\$0]					
FEE CALCULATION (continued)					
3. ADDITIONAL FEES					
Large Entity Fee Code (\$)	Small Entity Fee Code (\$)	Fee Description			Fee Paid
105	130	205	65	Surcharge - late filing fee or oath	
127	50	227	25	Surcharge - late provisional filing fee or cover sheet	
139	130	239	65	Non-English specification	
147	2,520	147	2,520	For filing a request for reexamination	
112	920	112	920	Requesting publication of SIR prior to Examiner action	
113	1,790	113	1,790	Requesting publication of SIR after Examiner action	
115	110	215	55	Extension for response within first month	
116	400	216	200	Extension for response within second month	
117	950	217	475	Extension for response within third month	
118	1,510	218	755	Extension for response within the fourth month	
128	2,060	228	1,030	Extension for response within the fifth month	
119	310	219	155	Notice of Appeal	
120	310	220	155	Filing a brief in support of an appeal	
121	270	221	135	Request for oral hearing	
140	110	240	55	Petition to revive unavoidably abandoned application	
141	1,320	241	660	Petition to revive unintentionally abandoned application	
142	1,320	242	660	Utility issue fee (or reissue)	
143	450	243	225	Design issue fee	
144	670	244	335	Plant issue fee	
122	130	122	130	Petitions to the Commissioner	
123	50	123	50	Petitions related to provisional applications	
126	240	126	240	Submission of Information Disclosure Statement	
581	40	581	40	Recording each patent assignment per property (times number of properties)	
149	790	249	790	For each additional invention to be examined (37 CFR 1.129(b))	
Other fee (specify)					SUBTOTAL (3) \$0
* Reduced by Basic Filing Fee Paid					

FEE TRANSMITTAL
PAGE 2

Complete If Known	
Application Number	
Filing Date	
First Named Inventor	Johannsson
Group Art Unit	
Examiner Name	

SUBMITTED BY		Complete (if applicable)		
Typed or Printed Name	Burton A. Amernick	Registration Number	24,852	
Signature		Date	3-31-98	Deposit Account User ID

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

APPLICANT: Johannsson et al
SERIAL NO: To be assigned GROUP NO: To be assigned
FILING DATE: Concurrently herewith EXAMINER: To be assigned
FOR: Use of Growth
Hormone

PRELIMINARY AMENDMENT

Honorable Commissioner of Patents and Trademarks
Washington, D.C. 20231

Sir:

Prior to initial examination, please amend the above-captioned case as follows.

In the Claims:

Claim 3, line 1, delete "or 2".

Claim 4, line 1, change "any preceding claims" to -claim 1-.

Claim 5, change "any preceding claims" to -claim 1-.

Claim 6, change "any preceding claims" to -claim 1-.

Claim 7, change "any preceding claims" to -claim 1-.

Claim 8, change "any preceding claims" to -claim 1-.

Claim 9, change "any preceding claims" to -claim 1-.

Claim 10, change "any preceding claims" to -claim 1-.

Please add the following new claims.

11. Use according to claim 2 in which the medicament is for treatment of abdominal/visceral obesity.
12. Use according to claim 2 in which the medicament is for increase insulin sensitivity.
13. Use according to claim 2 in which the medicament is for decrease of insulin resistance.
14. Use according to claim 2 in which the medicament is for decrease of total body fat mass.
15. Use according to claim 2 in which the medicament is for decrease of total body fat in human with abdominal/visceral obesity.
16. Use according to claim 2 in which the medicament is for visceral adipose tissue area.
17. Use according to claim 2 in which the medicament is for decrease of serum concentration of triglyceride.
18. Use according to claim 2 in which the medicament is for decrease of diastolic blood pressure.

Remarks

The claims have been amended to eliminate multiple dependency and to improve their format. None of these amendments is believed to involve any new matter. Accordingly, it is respectfully requested that the foregoing amendments be entered, that the application as so

amended receive an examination on the merits, and that the claims as now presented receive an early allowance.

Respectfully submitted,



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USE OF GROWTH HORMONE

The present invention relates to the use of growth hormone, preferably human
5 growth hormone or analogues thereof for the manufacturing of a medicament for
treatment of individuals with the Metabolic syndrome (also labeled Syndrome X).
They include individuals with abdominal/visceral obesity and its metabolic and
circulatory consequences including insulin resistance, lipoprotein aberrations and
hypertension. The medicament is also used to increase insulin sensitivity and for
10 treatment and prevention of non-insulin dependent diabetes mellitus.

Introduction

Striking similarities exhibits between The Metabolic Syndrome ^{1, 2} (also labeled
Syndrome X or Primary Insulin Resistance Syndrome) and untreated GH
15 deficiency in adults ³. The most central findings in both these syndromes are
abdominal/visceral obesity and insulin resistance ^{1, 4-6}. Other features common
to both conditions are high triglyceride and low high-density lipoprotein
cholesterol concentrations, an increased prevalence of hypertension, elevated
levels of plasma fibrinogen and plasminogen activator inhibitor-1 activity,
20 premature atherosclerosis and increased mortality from cardiovascular diseases ^{1,}
^{4, 7-11}.

The Metabolic Syndrome is associated with multiple endocrine abnormalities.
They include increased cortisol secretion, blunted secretion of gonadotrophins and
sex steroids and abnormalities in the GH/insulin-like growth factor-I (IGF-I) axis
25 12-14. With increased adiposity GH secretion is blunted with decreased mass of

GH secreted per burst but without any major impact on GH secretory burst frequency ¹⁵. The serum IGF-I concentration is principally GH dependent and influences GH secretion through a negative feed-back system ¹⁶. The serum levels of IGF-I are inversely related to the percentage body fat ¹⁵. In addition, we have
5 previously shown that the low serum IGF-I concentration in obesity is predominantly related to the amount of visceral adipose tissue and not to the amount of subcutaneous fat mass ¹³. These findings, together with other endocrine disturbances in central obesity, suggest that the low GH secretion which is observed is secondary to a central disturbance of the neuroendocrine regulation,
10 including the GH/IGF-I axis.

The abdominal/visceral obesity and insulin resistance observed in The Metabolic Syndrome constitute the base for hypertension, dyslipoproteinemia and non-insulin dependent diabetes mellitus.¹

Replacement therapy with recombinant human GH (rhGH) has demonstrated favorable effects on most of the features of GH deficiency in adults ¹⁷. Whether rhGH treatment can improve the metabolic abnormalities observed in abdominal/visceral obesity has never been investigated. In the present study a randomized, double-blind, placebo-controlled design was used to evaluate the effects of rhGH administration in patients with abdominal/visceral obesity
15

20 In the present study thirty men, 48 to 66 years of age with abdominal/visceral obesity were treated with recombinant human GH (rhGH) in a 9-month randomized, double-blind, placebo-controlled trial. Body fat was assessed from total body potassium and abdominal subcutaneous and visceral adipose tissue was measured using computed tomography. Glucose disposal rate (GDR) was
25 measured during euglycemic, hyperinsulinemic glucose clamp.

In response to the rhGH treatment, total body fat, abdominal subcutaneous and visceral adipose tissue decreased. The GDR (glucose disposal rate) increased in the rhGH-treated group as compared with the placebo-treated one. The mean serum concentrations of total cholesterol and triglyceride decreased while blood glucose and serum insulin concentrations were unaffected by the rhGH treatment.
5 Furthermore, diastolic blood pressure decreased and systolic blood pressure was unchanged in response to rhGH treatment.

The present study has demonstrated that GH can favorably affect some of the multiple perturbations associated with The Metabolic Syndrome. This includes a
10 reduction in abdominal/visceral obesity an improved insulin sensitivity and favorable effects on lipoprotein metabolism and diastolic blood pressure. Thus, the prevention and treatment of non-insulin dependent diabetes mellitus may be possible.

The findings must be regarded as unexpected and surprising and of utmost interest giving a possibility to treat and prevent diseases associated with increased
15 cardiovascular morbidity and mortality.

The present invention thus relates to the use of growth hormone or analogues thereof as claimed in the claims on file.

By analog is meant a substance having the same biological activity as described
20 here and having at least 65 % homology with natural occurring growth hormone.

Legends to figures.

Figure 1. Mean total body fat calculated from total body potassium, abdominal subcutaneous adipose tissue (AT) area at the level of L4-L5 and volume of visceral
25 AT.

Figure 2. Abdominal subcutaneous and visceral adipose tissue determined with computed tomography at the level of L4-5 in one man before (A) and after 9 months of rhGH treatment (B).

Figure 3. Mean fasting blood glucose, serum insulin and glucose disappearance

5 rate (GDR).

EXAMPLE

Subjects and methods

Patients

10 Thirty men (48 to 66 years of age) were studied (Table 1). They were recruited by advertisement in a local newspaper. The criteria for inclusion in the study were age approximately 50 to 65 years with a body mass index between 25 and 35 kg/m², an IGF-I less than 160 µg/L (low normal)¹⁸ and a waist hip ratio of more than 0.95. The criteria for exclusion were overt diabetes mellitus, previous
15 cardiovascular event or heart disease.

In the rhGH treated group, two patients were receiving treatment for hypertension with both atenolol (100 mg per day) and nifedipin (40 mg per day) and one patient had slight asthma, treated with salmeterol and terbutalin inhalations. In the placebo treated group, one patient had a slight depressive disorder and was receiving paroxetin (10 mg per day). These medications were kept stable during
20 the study period.

Study design

This study was a 9 month randomized, double-blind, placebo-controlled trial of the administration of rhGH. Informed consent was obtained from each patient

before the study. The study was approved by the Ethics Committee at the University of Göteborg and by the Swedish Medical Products Agency, Uppsala, Sweden.

Treatment

5 The daily rhGH dose was 9.5 µg/kg (0.20 IU/kg body weight/week), administrated subcutaneously before bedtime. The dose was reduced by half in the event of side-effects. The average dose reduction during the 9-month study was 0.17 mg per day (range -1.7 to 0). The placebo vials contained the same vehicle as the rhGH vials and both preparations were visually indistinguishable.

10 Compliance assessed by counting the returned empty vials and expressing that number as a percentage of vials needed for the treatment period was 87.3 % (range 32-100).

Study protocol

15 The patients were studied as out-patients before and after 6 weeks, 6 and 9 months of treatment. Physical and laboratory examinations were performed at all visits. The euglycemic hyperinsulinemic clamp, adipose tissue needle aspiration and measurement of blood pressure and heart rate were performed at baseline, 6 weeks and after 9 months of treatment. Body weight was measured in the morning to the nearest 0.1 kg wearing indoor clothing, and body height was measured

20 barefoot to the nearest 0.01 m. The body mass index was calculated as body weight in kilograms divided by height in meters squared. Waist circumference was measured in the standing position with a flexible plastic tape midway between the lower rib margin and the iliac crest, and the hip girth at the widest part of the hip. Systolic and diastolic blood pressure were measured after 5 minutes of supine rest

25 using the sphygmomanometric cuff method. This measurement was repeated after one minute and the mean value was used.

Total-body potassium was measured by counting the emission of 1.46 MeV gamma-radiation from the naturally occurring ^{40}K isotope in a high-sensitive 3π whole body counter with a coefficient of variation (CV) of 2.2%. Fat-free mass (FFM) was estimated by assuming a potassium content of 64.7 mmol/kg FFM¹⁹

5 and body fat (BF) was calculated as body weight-FFM. A five-scan computed tomography technique was used (Philips Tomoscan 350) to measure abdominal adipose tissue. The five levels were derived from two scanograms and included one scan at the level of the mid-thigh with the lower edge of symphysis as a reference point. The other four levels were the lower edge of symphysis, L4-5

10 lumbar disc, L3-4 lumbar disc and a scan at the level of liver and spleen. The tissue areas and anatomic boundaries were determined as described previously²⁰. Total volume of visceral adipose tissue was determined from the five-scan model. Sagittal diameter and abdominal subcutaneous and visceral adipose tissue areas were determined at the level of L4-5.

15 A euglycemic, hyperinsulinaemic glucose clamp was performed after an overnight fast as previously described¹⁴. In brief, insulin was infused together with glucose into a venous catheter with the tip at the level of the axillary vein at appropriate rates to obtain submaximal insulin concentrations. Blood glucose concentrations were monitored every 10 min. and glucose infusion rate was adjusted to fasting

20 levels. Glucose disappearance rate (GDR) was measured for 20 minutes in steady-state conditions, which was reached after 100 min. The insulin concentrations during steady state were $214 \pm 10 \mu\text{IU}/\text{mL}$ before treatment, $226 \pm 12 \mu\text{IU}/\text{mL}$ at 6 weeks and $213 \pm 11 \mu\text{IU}/\text{mL}$ at 9 months. During the clamp, in steady-state conditions, a subcutaneous abdominal adipose tissue biopsy was obtained by

25 needle aspiration for the determination of lipoprotein lipase (LPL) activity. The needle aspiration was performed under local anesthesia, 0.1 m lateral to the umbilicus. The biopsies were immediately frozen in liquid nitrogen and stored at -80-C until assay.

Analytic methods

Total LPL activity in adipose tissue was determined after homogenization of the tissue in the detergent-containing buffer as described previously 21. Bovine skim milk was used as a standard to correct for interassay variation. The amount of

5 triglycerides (TG) in the tissue was measured after extraction 22 and weighed after evaporation of solvents. Activity was expressed in milliunits (1mU=1 mmol FFA released per min.) per gram adipose tissue and per gram TG. Control experiments showed that the assay was linear with the amount of sample and incubation time over the range used. The within-assay CV was 4.3%.

10 Blood samples were drawn in the morning after an overnight fast. The serum concentration of insulin-like growth factor-I (IGF-I) was determined by a hydrochloric acid-ethanol extraction radioimmunoassay using authentic IGF-I for labeling (Nichols Institute Diagnostics, San Juan Capistrano, CA, USA) with within-assay CV of 2.5% and 4.2% at serum concentrations of 125 μ g/L and 15 345 μ g/L respectively. The standard deviation score for IGF-I was then calculated from predicted IGF-I values adjusted for age and sex obtained from the normal population 18.

The insulin-like growth factor binding protein-3 (IGFBP-3) concentration in serum was determined by radioimmunoassay (Nichols Institute Diagnostics) with total 20 CV of 6.2% and 5.7% at serum concentrations of 2.05 mg/L and 3.49 mg/L respectively.

Serum cholesterol and triglyceride concentrations were determined with enzymatic methods (Boehringer, Mannheim, Germany). The within-assay CV for total cholesterol and triglyceride determinations was 0.9% and 1.1% respectively.

25 Fibrinogen was measured according to a syneresis method 23 with a total CV of 4% at 2.5 g/L. Plasminogen activator inhibitor (PAI)-1 activity was measured

using a Spectrolyse (pL) PAI kit (Biopool Stablyte, Umeå, Sweden) with a total CV of 10% at concentrations between 10-40 IU/ml.

Serum insulin was determined by a radioimmunoassay (Phadebas, Pharmacia, Uppsala, Sweden) and blood glucose was measured by the glucose-6-phosphate dehydrogenase method (Kebo Lab, Stockholm, Sweden). Hemoglobin A1c (HbA1c) was determined by high pressure liquid chromatography (Waters, Millipore AB, Sweden) and C-peptide was determined by a radioimmunoassay (Santec, USA). Free fatty acids (FFA) levels were determined using an enzymatic colorimetric method (NEFAC; Wako, Neuss, Germany), while sex hormone binding globulin (SHBG) was measured by an immunoradiometric assay (Farmos, Diagnostica, Åbo, Finland) and total testosterone by a radioimmunoassay (ICM, Biomedical, Costa Mesa, USA).

Statistical methods

All descriptive statistical results are presented as the means and standard error of the mean (SEM). Comparisons between the two treatment groups were performed by a two-way analysis of variance (ANOVA) for repeated measurements. Comparisons between baseline values and values at 9 months in the two groups were performed with the Students' t-test for independent groups. Correlations were sought by calculating the Pearson linear correlation coefficient. Pearson's chi square test was used to test the independence of frequency of hypertension and smoking between the rhGH and placebo treated groups. Before statistical analysis, logarithmic transformation of data with skewed distribution was performed. A two-tailed probability value of less than 0.05 was considered significant.

Results

The two groups were matched with regard to age, body height, body weight and waist hip ratio, and did not differ significantly regarding number of subjects with

medically treated hypertension (Chi χ^2 =1.87; $p=0.17$) and current smokers (Chi χ^2 =1.2; $p=0.27$) (Table 1). At baseline the two groups did not differ significantly from each other in any of the variables studied.

Body composition

5 The patients' mean overall body mass index and fat-free mass did not change while mean total body fat decreased by $9.2 \pm 2.4\%$ during rhGH treatment in comparison with placebo treatment (Figure 1). Waist circumference and sagittal diameter decreased in response to rhGH while no change occurred in the placebo group. Moreover, abdominal subcutaneous and visceral adipose tissue area at the 10 level of L4-5 decreased in response to rhGH by $6.1 \pm 3.2\%$ and $18.1 \pm 7.6\%$ respectively (Figure 1 and Table 2). The corresponding values in the placebo group were $+2.0 \pm 2.8\%$ and $-3.2 \pm 7.6\%$ respectively. Moreover, the volume of visceral adipose tissue decreased in the rhGH treated group by $17.9 \pm 3.5\%$ while no change was observed in the placebo treated group ($-0.2 \pm 4.2\%$) (Figure 1).

15 Thus, the percentage of visceral adipose tissue of the total adipose tissue at the level of L4-5 decreased by $14.5 \pm 3.8\%$, while the percentage of subcutaneous adipose tissue of the total adipose tissue at the level of L4-5 increased by $5.4 \pm 1.7\%$ in the rhGH-treated group.

Abdominal subcutaneous and visceral adipose tissue was determined with 20 computed tomography at the level of L4-5 in one man before (A) and after 9 months of rhGH treatment (B). The scan showed a reduction of both visceral and subcutaneous adipose tissue . (Figure 2)

Glucose metabolism

No significant effects were elicited by rhGH treatment on blood glucose, serum 25 insulin and HbA1c (Table 3 and Figure 3). The serum concentration of C-peptide

increased in the rhGH treated group due to a transient increase at 6 weeks, but the C-peptide level was similar in the two groups at 9 month.

In the rhGH treated group the GDR showed an initial decrease after 6 weeks of treatment followed by an increment while the placebo-treated group
 5 demonstrated a slight reduction over time (Figure 3). The average increase in GDR in response to 9 months of rhGH was 1.2 ± 0.7 mg/kg/min compared with baseline while in the group receiving placebo a slight decrease by 0.4 ± 0.6 mg/kg/min occurred. This difference between the two groups was still found after correcting the GDR for the amount of fat-free mass.
 10 In the rhGH treated group, an inverse correlation was found between the change in GDR and change in serum triglyceride concentration ($r=-0.58$; $p<0.05$) but no significant correlations were found between the change in GDR and changes in serum IGF-I concentration ($r=-0.24$), serum insulin concentration ($r=-0.27$), LPL activity ($r=0.09$), diastolic blood pressure ($r=-0.30$), total body fat ($r=-0.06$) and volume of visceral adipose tissue ($r=-0.11$).
 15

Total cholesterol, triglycerides, LPL activity, plasma fibrinogen and PAI-1 activity
 (Table 3)

The mean total cholesterol concentration decreased from 6.1 ± 0.2 to 5.4 ± 0.3 mmol/L and the triglyceride concentration decreased from 2.09 ± 0.29 to 1.78 ± 0.23 mmol/L in response to rhGH treatment. The corresponding changes in the placebo group were from 5.4 ± 0.3 to 5.5 ± 0.2 mmol/L and from 1.65 ± 0.13 to 2.05 ± 0.26 mmol/L respectively.
 20

The mean total LPL activity in subcutaneous abdominal tissue did not change during rhGH treatment in comparison with placebo treatment. However, between 25 6 weeks and 9 months, LPL activity tended to increase in the rhGH treated group

as compared with the placebo treated group ($p=0.06$). Similar results were obtained when LPL activity was expressed in mU/g TG.

The plasma concentration of fibrinogen increased in response to rhGH while PAI-I activity was unaffected by rhGH treatment as compared with placebo.

5 Blood pressure and heart rate

Diastolic blood pressure decreased from 75 ± 2 to 70 ± 2 mmHg in the rhGH treated group. The corresponding values in the placebo group were from 73 ± 2 to 74 ± 2 mmHg ($p<0.05$). No significant effects on systolic blood pressure or heart rate were observed between the two treatment groups.

10 Serum IGF-I, IGFBP-3, testosterone and SHBG (Table 4)

Before treatment, the serum IGF-I concentration was low normal in both treatment groups. A significant increment in serum IGF-I and IGFBP-3 concentrations occurred in the rhGH treated group as compared with placebo treatment. At 6 weeks, the serum IGF-I concentration reached on average 3.30 ± 0.35 SD above the predicted mean in the rhGH treated group while at 9 month the mean serum IGF-I concentration was 1.89 ± 0.48 SD above the predicted mean. Serum concentrations of testosterone and SHBG were not significantly affected by rhGH treatment.

15 Side-effects

No drop-outs occurred during the study. Side-effects were observed in 8 subjects in the rhGH treated group, and were mainly a result of fluid retention. Five had peripheral edema, two subjects experienced muscle stiffness and arthralgia, one developed mild carpal tunnel syndrome and one subject experienced increased perspiration. These side-effects appeared during the first 6 weeks of treatment and subsided in four patients in response to a reduction in dose performed within 4 months after start of treatment; in three patients, the side-effects subsided

spontaneously. One man from the rhGH treated group with medically treated hypertension was taken off treatment after 8 months when he experienced an intracerebral hemorrhage. Three subjects in the placebo group experienced slight and transient peripheral edema.

5 **Legends to figures.**

Figure 1.

Mean total body fat calculated from total body potassium, abdominal subcutaneous adipose tissue (AT) area at the level of L4-L5 and volume of visceral AT assessed with computerized tomography during 9 months of treatment with rhGH or placebo in 30 men with abdominal/visceral obesity. The horizontal bars indicate the SE for the mean values shown and p-values denotes the differences between the two groups by two-way ANOVA for repeated measurements.

Figure 2.

Abdominal subcutaneous and visceral adipose tissue determined with computed tomography at the level of L4-5 in one man before (A) and after 9 months of rhGH treatment (B). The scan shows the reduction of both visceral and subcutaneous adipose tissue (shown as dark gray area in the figure).

Figure 3.

Mean fasting blood glucose, serum insulin and glucose disappearance rate (GDR) assessed with euglycaemic hyperinsulinaemic glucose clamp during 9 months of treatment with rhGH or placebo in 30 men with abdominal/visceral obesity. The horizontal bars indicate the SE for the mean values shown and p-values denotes the differences between the two groups by two-way ANOVA for repeated measurements.

Discussion

We have shown that 9 months of rhGH treatment in middle-aged men with abdominal/visceral obesity reduced total body fat and resulted in a specific and marked decrease of both abdominal subcutaneous and visceral adipose tissue.

5 Insulin sensitivity assessed with euglycemic glucose clamp technique improved and serum concentrations of total cholesterol and triglyceride decreased. Furthermore, diastolic blood pressure decreased.

The men who were studied were moderately obese with a preponderance of abdominal and/or visceral localization of body fat as judged from comparisons 10 with randomly selected men of comparable age from the same city²⁴. As a group, they had slight to moderate metabolic changes known to be associated with abdominal/visceral obesity with moderate insulin resistant as judged from the GDR values obtained during the euglycemic glucose clamp, although no one had overt diabetes.

15 Although, we used a lower daily rhGH dose than previously reported in trials studying healthy adults^{25, 26}, the initial rhGH doses administrated were apparently too high as judged by the frequency of side-effects and initial high average serum IGF-I concentrations. After dose reduction the average serum IGF-I concentration where within the normal range indicating that the doses of rhGH 20 during the latter part of the study were more physiological. This might in turn explain the less marked anabolic action of the rhGH treatment demonstrated in this study in comparison with previous trials in healthy adults^{25, 26}.

The marked effect of GH replacement on body composition in GH-deficient adults has been a consistent observation in many studies¹⁷. The profound lipolytic effect 25 of GH was also demonstrated in the present study with a preferential reduction in visceral adipose tissue depots²⁷. These changes have been associated with a GH-

induced reduction in the antilipolytic effects of insulin, which is markedly different in different adipose tissue regions ²⁸.

GH exerts direct insulin-antagonistic effects even after the administration of physiological doses of rhGH. GH has been considered to be the principal factor in
5 the decrease in insulin sensitivity observed in the early morning, the so called "dawn phenomenon" ²⁹ and the insulin resistance following hypoglycemia ³⁰.

Thus, our observation of increased insulin sensitivity during prolonged rhGH treatment is unexpected although not inexplicable. We have previously shown that 6 weeks of rhGH treatment in GH-deficient adults only induced a temporary decrease in insulin sensitivity which after 6 months of treatment was restored to baseline values ³¹. The response of GDR to rhGH treatment on in this trial showed a similar initial tendency to induce insulin resistance but after 9 months a marked improvement was found. This improvement could be explained by the decreased visceral adipose tissue mass induced by GH, followed by decreased FFA exposure to the liver counteracting the insulin-antagonistic effects of GH. Alternatively, as the major site of glucose disposal is in the skeletal muscle ³² it has to be considered that the improvement in GDR in response to the more prolonged rhGH treatment might also be an effect of increased glucose transport in the skeletal muscle possibly mediated through the IGF-I receptor ³³, increased proportion of insulin-sensitive type I muscle fibers ³⁴ or increased capillarization in the skeletal muscle.
20

GH has been found to be an important regulator of the hepatic LDL-receptor ³⁵ and the overall lipoprotein metabolism ³⁶. The reduction in total cholesterol is conceivable an effect of enhanced hepatic LDL receptor activity in response to GH³⁵. In healthy adults, short-term rhGH administration has been reported to
25 increase serum triglyceride concentration ²⁵. In this study, the serum triglyceride concentration also displayed an initial increase in response to rhGH treatment. This could be an effect of both an increased flux of FFA to the liver and a direct

stimulatory effect on the esterification of oleic acid into triglyceride and phospholipids in hepatocytes³⁷ in response to GH which in turn enhances the very low density lipoprotein production from the liver. However, after 9 months of rhGH treatment serum triglyceride concentration had decreased again,
5 probably as an effect of the increased insulin-stimulated glucose uptake which is known to be inversely related with the very low density lipoprotein secretion rate from the liver and serum triglyceride levels⁴.

The initial tendency of a decline in LPL activity observed in the present study is in accordance with a previous 2 weeks treatment trials with rhGH³⁸. During the
10 more prolonged rhGH treatment, however, LPL activity tended to increase which may, at least in part, explain the decrease in serum triglyceride concentration at 9 months. The changes in GDR and LPL activity showed a similar biphasic pattern in response to GH. It may thus be speculated that the suggested influence of GH on LPL activity is mediated through insulin sensitivity, since insulin is known to
15 be a potent stimulator of LPL.

Nine months of rhGH treatment reduced diastolic blood pressure without affecting systolic blood pressure. This is in line with results from GH-deficient adults where rhGH administration reduced diastolic blood pressure possibly as an
20 effect of reduced peripheral vascular resistance³⁹. The mechanisms behind the reduction in peripheral vascular resistance might be indirect through the reduced abdominal obesity and increased insulin sensitivity⁴⁰ or more direct through the action of IGF-I on the vascular wall⁴¹.

Abdominal/visceral obesity is associated with increased plasma fibrinogen concentration and PAI-1 activity^{42, 43} which both are established risk factors for
25 myocardial infarction and stroke^{44, 45}. The slightly increased plasma fibrinogen concentration in response to GH may be mediated through increased serum IGF-I concentration¹⁸. In GH-deficient adults, 2 years of rhGH treatment tended to

decrease plasma fibrinogen levels and diminish PAI-1 activity⁴⁶. This further illustrates the importance of the duration of rhGH treatment on the metabolic effects of GH.

Previous studies have shown that testosterone treatment of middle-aged men with abdominal/visceral obesity induced improved insulin sensitivity, plasma lipid levels, diastolic blood pressure as well as a specific decrease of visceral adipose tissue mass^{14, 46}. Since testosterone treatment of men with hypogonadotropic hypogonadism increases GH secretion⁴⁷ the similarities between testosterone and rhGH treatment might be explained by increased GH levels or by additive or synergistic effects of GH and testosterone on adipose tissue metabolism⁴⁸.

The multiple endocrine alterations associated with abdominal/visceral obesity can either be primary responsible or be the consequence of the obese condition. This is the first trial clearly to demonstrate favorable effects of GH on the multiple perturbations associated with abdominal/visceral obesity. We therefore suggest that a blunted GH secretion could be an important factor in the development of the metabolic and circulatory consequences of abdominal/visceral obesity. The metabolic effects demonstrated in this study are probably of importance for the risk of cardiovascular morbidity and mortality in men with abdominal and central adiposity.

Table 1.

Clinical characteristics of the cohort of 30 men treated with GH or placebo for 9 months.

Patient characteristics	GH	Placebo
Number of men	16	14
Mean age (range); years	58.3 (48-66)	57.9 (52-63)
Treated hypertension; no.	2	0
Present smokers; no.	5	2
Body height; m	$1.80 \pm 0.02^*$	1.77 ± 0.02
Body weight; kg	101.7 ± 2.4	96.2 ± 2.6
Waist hip ratio	1.01 ± 0.01	1.03 ± 0.02

* Plus-minus values are means \pm standard error of the mean.

Table 2.

Measurements of body mass index (BMI), fat free mass, waist circumference, sagittal diameter and abdominal visceral adipose tissue (AT) area at the level of L4-5 in 30 men, during 9 months of rhGH or placebo treatment.

Variable	Baseline	6 weeks	6 months	9 months	P #
BMI; kg/m²					
GH	31.4±0.7	31.6±0.7	31.3±0.7	31.1±0.8	0.2
Placebo	30.5±0.8	30.6±0.7	30.5±0.8	30.7±0.8	
Fat free mass; kg					
GH	67.5±1.8	69.7±1.9	71.1±1.8	69.5±2.2	0.4
Placebo	64.6±1.4	65.2±1.4	66.4±1.3	65.3±1.4	
Waist; cm					
GH	111.8±1.8	110.8±1.8	107.6±1.7	109.8±1.9	0.002
Placebo	109.5±2.5	109.4±2.4	109.3±2.3	111.0±2.3	
Sagittal diameter; cm					
GH	26.1±0.5	25.9±0.5	25.2±0.5	25.0±0.6	0.03
Placebo	25.5±0.7	25.3±0.8	24.6±0.9	25.5±0.8	
Visceral AT; cm²					
GH	126±15	121±19	98±14	99±15	0.004
Placebo	163±16	147±13	142±12	150±13	

All values are expressed as the mean (SEM).

P-values denote the differences between the two groups by two-way ANOVA for repeated measurements.

Table 3.

Measurements of glycosylated hemoglobin (HbA1c), C-peptide, total cholesterol, triglycerides, lipoprotein lipase (LPL) activity expressed as mU per gram of adipose tissue (AT), plasma fibrinogen and plasminogen activator inhibitor (PAI)-1 in 30 men, during 9 months of rhGH or placebo treatment.

Variable	Baseline	6 weeks	6 months	9 months	P #
HbA1c; %					
GH	5.2±0.1	5.2±0.1	5.4±0.1	5.3±0.1	0.8
Placebo	4.9±0.1	4.9±0.1	5.0±0.1	5.1±0.2	
C-peptide; ng/L					
GH	1.11±0.10	1.60±0.13	1.30±0.17	1.18±0.07	0.02
Placebo	1.28±0.27	1.30±0.27	1.28±0.32	1.42±0.30	
Total cholesterol; mmol/L					
GH	6.1±0.2	5.7±0.2	6.1±0.2	5.4±0.3	0.006
Placebo	5.4±0.3	5.6±0.2	6.1±0.3	5.5±0.2	
Triglycerides; mmol/L					
GH	2.09±0.29	2.60±0.42	2.31±0.23	1.78±0.23	0.02
Placebo	1.65±0.13	1.80±0.20	1.85±0.22	2.05±0.26	
LPL activity; mU/g AT					
GH	287±26	205±15	ND	371±65	0.2
Placebo	224±31	228±29	ND	263±25	
Fibrinogen; g/L					
GH	3.13±0.13	3.44±0.10	3.49±0.13	3.24±0.19	0.04
Placebo	3.03±0.11	2.73±0.09	3.39±0.28	2.84±0.10	
PAI-1 activity; U/mL					
GH	31.3±6.4	23.2±2.6	18.3±3.1	22.9±3.5	0.4
Placebo	17.0±3.8	22.9±4.2	19.9±4.5	27.1±6.3	

All values are expressed as the mean (SEM).

P-values denote the differences between the two groups by two-way ANOVA for repeated measurements.

Table 4.

Measurements of serum IGF-I, IGFBP-3, free fatty acids (FFA), testosterone and steroid hormone binding globulin (SHBG) in 30 men, during 9 months of GH/placebo treatment.

Variable	Baseline	6 weeks	6 months	9 months	P #
IGF-I; µg/L					
GH	134±8	338±16	320±23	268±23	<0.001
Placebo	120±11	121±11	129±12	119±12	
IGF-I SD score *					
GH	-0.82±0.17	3.30±0.35	2.91±0.49	1.89±0.48	<0.001
Placebo	-1.07±0.23	-1.04±0.23	-0.87±0.24	-1.08±0.25	
IGFBP-3; mg/L					
GH	2.36±0.13	3.20±0.10	3.19±0.13	2.71±0.14	0.001
Placebo	2.10±0.16	2.21±0.17	2.45±0.21	2.18±0.19	
FFA; µmol/L					
GH	0.77±0.07	0.99±0.11	0.78±0.11	0.75±0.07	0.5
Placebo	0.73±0.05	0.73±0.05	0.59±0.06	0.67±0.05	
Testosterone; nmol/L					
GH	14.6±1.2	12.9±1.0	13.9±1.3	12.4±1.1	0.12
Placebo	14.6±1.0	15.4±1.2	17.6±2.1	14.6±1.3	
SHBG; nmol/L					
GH	26.4±3.4	25.5±3.5	25.9±3.1	23.4±2.8	0.8
Placebo	28.1±2.9	29.1±3.1	29.2±3.2	25.5±2.7	

All values are expressed as the mean (SEM).

P-values denotes the differences between the two groups by two-way ANOVA for repeated measurements.

* The standard deviation score for serum IGF-I is calculated from predicted IGF-I values adjusted for age and sex obtained from the normal population ¹⁸.

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CLAIMS

1. Use of growth hormone, preferably human growth hormone, or analogues thereof for the manufacturing of a medicament for treatment of The Metabolic

5 Syndrome.

2. Use of growth hormone, preferably human growth hormone, or analogues thereof for the manufacturing of a medicament for prevention and treatment of non-insulin dependent diabetes mellitus.

10 3. Use according to claim 1 or 2 in which the medicament is for treatment of abdominal/visceral obesity.

4. Use according to any preceding claims in which the medicament is for increase of insulin sensitivity.

5. Use according to any preceding claims in which the medicament is for decrease of insulin resistance.

15 6. Use according to any preceding claims in which the medicament is for decrease of total body fat mass.

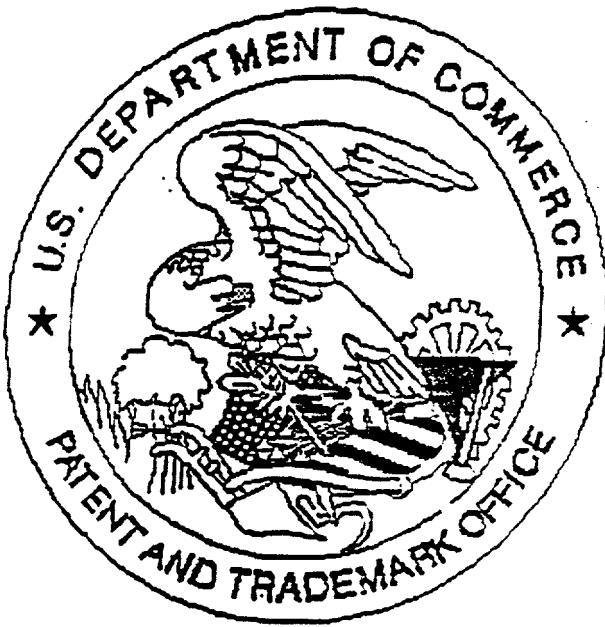
7. Use according to any preceding claims in which the medicament is for decrease of total body fat in human with abdominal/visceral obesity.

20 8. Use according to any preceding claims in which the medicament is for visceral adipose tissue area.

9. Use according to any preceding claims in which the medicament is for decrease of serum concentration of triglyceride.

10. Use according to any preceding claims in which the medicament is for decrease of diastolic blood pressure.

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